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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/560,774	12/14/2005	Fabien Schweighoffer	BJS-3665-167	5165
23117 NIXON & VAN	7590 10/14/200 NDERHYE, PC	EXAMINER		
901 NORTH G	LEBE ROAD, 11TH F	JAVANMARD, SAHAR		
ARLINGTON, VA 22203			ART UNIT	PAPER NUMBER
			1627	
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			10/14/2009	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)	
	10/560,774	SCHWEIGHOFFER ET AL.	
Office Action Summary	Examiner	Art Unit	
	SAHAR JAVANMARD	1627	
The MAILING DATE of this communication ap Period for Reply	ppears on the cover sheet with the o	correspondence address	
A SHORTENED STATUTORY PERIOD FOR REP WHICHEVER IS LONGER, FROM THE MAILING I - Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory perio- Failure to reply within the set or extended period for reply will, by statu Any reply received by the Office later than three months after the mail earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICATION 1.136(a). In no event, however, may a reply be tind will apply and will expire SIX (6) MONTHS from the, cause the application to become ABANDONE	N. mely filed the mailing date of this communication. ED (35 U.S.C. § 133).	
Status			
Responsive to communication(s) filed on 31.  2a) This action is <b>FINAL</b> . 2b) Th  3) Since this application is in condition for allow closed in accordance with the practice under	is action is non-final. ance except for formal matters, pro		
Disposition of Claims			
4)  Claim(s) 12-14 is/are pending in the applicati 4a) Of the above claim(s) is/are withdrest is/are allowed.  5)  Claim(s) is/are allowed.  6)  Claim(s) 12-14 is/are rejected.  7)  Claim(s) is/are objected to.  8)  Claim(s) are subject to restriction and/  Application Papers  9)  The specification is objected to by the Examin	awn from consideration.  /or election requirement.		
10) The drawing(s) filed on is/are: a) according a deposition of the drawing not request that any objection to the Replacement drawing sheet(s) including the correct should be correctly as the deposition of the should be deposited to by the Equation is objected to by the Equation is objected to by the Equation is objected to by the Equation is objected.	ccepted or b) objected to by the e drawing(s) be held in abeyance. Se ection is required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority document application from the International Burest * See the attached detailed Office action for a list	nts have been received. nts have been received in Applicat iority documents have been receive au (PCT Rule 17.2(a)).	ion No ed in this National Stage	
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO/SB/08)  Paper No(s)/Mail Date 7/31/09.	4)  Interview Summary Paper No(s)/Mail D 5)  Notice of Informal F 6)  Other:	ate	

### **DETAILED ACTION**

# Status of the Application

This Office Action is in response to applicant's arguments filed on July 31, 2009. Claim(s) 12-14 are pending and are examined herein.

## Response to Arguments

Applicant's arguments with respect to the 103(a) rejection of claims 7 and 11-14 as being unpatentable over Ikhlef et al. (US Pub. No. 2003/0064374 A1) in view of Dalton et al. (WO 95/11887) has been fully considered and said rejection is hereby withdrawn.

In response to Applicant's contention that "the results presented in the present application are unexpected because, for example, they rely on a previously unknown mechanism of action of etazolate, not previously foreseen or predicted by the cited art, namely the modulation of GABAA receptor causing sAPPalpha production". In response to this argument, Examiner respectfully notes that "[T]he discovery of a previously unappreciated property of a prior art composition, or of a scientific explanation for the prior art's functioning, does not render the old composition patentably new to the discoverer." *Atlas Powder Co. v. Ireco Inc.*, 190 F.3d 1342, 1347, 51 USPQ2d 1943, 1947 (Fed. Cir. 1999). Thus the claiming of a new use, new function or unknown property which is inherently present in the prior art does not necessarily make the claim patentable. *In re Best*, 562 F.2d 1252, 1254, 195 USPQ 430, 433 (CCPA 1977).

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Furthermore, the claim recites a method of treatment, the mechanism by which it is achieved is not given patentable weight and is not a part of the search. As set forth on record, Ikhlef teaches the administration of etazolate in the treatment of Alzheimer's disease. Although the reference is silent on cognitive perception, one of ordinary skill in the art would expect with a reasonable degree of success to also try the regimen to treat the cognition aspect of Alzheimer's disease. In order to further justify that such an expectation is reasonable, a new rejection is set forth on record in the office action below.

### Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

Claims 12-14 are rejected under 35 U.S.C. 103(a) as being unpatentable over Ikhlef et al. (US Pub. No. 2003/0064374 A1) in view of Schumcher et al. (US Patent No. 7,153,871 B1).

Ikhlef teaches treating neurodegenerative diseases, including ALS and Alzheimer's disease with the use of etazolate, which is a PDE4 inhibitor (page 4, [0056]; claims 9, 12-14, and 17).

Ikhlef teaches that etazolate may be administered by any method known in the art preferably injection, namely the intravenous route (page 4, [0058]).

Ikhlef does not specifically teach treating cognitive deficits or the monitoring thereof.

Schumacher teaches a class of aminoindazole and aminobenzofuran analogs as a method of treatment that involves the inhibition of PDE4 enzymes. The invention includes methods of selective inhibition of PDE4 enzymes in animals, e.g., mammals, especially humans, wherein such inhibition has a therapeutic effect, such as where such inhibition may relieve conditions involving neurological syndromes, such as the loss of memory, especially long-term memory (column 2, lines 29-36).

Schumacher teaches the condition of memory impairment is manifested by impairment of the ability to learn new information and/or the inability to recall previously learned information. Memory impairment is a primary symptom of dementia and can also be a symptom associated with such diseases as Alzheimer's disease, schizophrenia, Parkinson's disease, Huntington's disease, Pick's disease,

Creutzfeld-Jakob disease, HIV, cardiovascular disease, and head trauma as well as age-related cognitive decline (column 2, lines 41-49).

Dementias are diseases that include memory loss and additional intellectual impairment separate from memory. The present invention includes methods for treating patients suffering from memory impairment in all forms of dementia.

Dementias are classified according to their cause and include: neurodegenerative dementias (e.g., Alzheimer's, Parkinson's disease, Huntington's disease, Pick's disease), vascular (e.g., infarcts, hemorrhage, cardiac disorders), mixed vascular and Alzheimer's, among others column 18, lines 50-55).

Schumacher teaches two methods of monitoring cognitive improvement upon administration of the PDE4 inhibitors, including *in vivo* testing for learning and memory 1)passive avoidance in rate, 2) radial arm maze task (column 26, example 11, methods A and B)

It would have been obvious to one of ordinary skill in the art at the time of the invention to have treated Alzheimer's disease with the administration of etazolate, as taught by Ikhlef and also treated perceptive cognition. The motivation, provided by Schumacher, teaches that PDE4 inhibition is employed as a method of treatment for neurodegenerative dementias, namely Alzheimer's disease, to improve loss of memory, especially long term memory. Thus, it would have been obvious to one of ordinary skill in the art to have expected, with a reasonable degree of success that a PDE4 inhibitor, namely etazolate, which is effective in treating Alzheimer's disease, can also be employed to improve cognition based on reasons of record. One would expect that

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because both classes of compounds are PDE4 inhibitors and employed to treat

Alzheimer's disease that one of ordinary skill in the art would expect, with a reasonable
degree of success, that the former, namely etazolate, would also be an obvious
candidate, to at least try, as a method of improving the cognitive aspect of Alzheimer's
disease. Additionally, it would have been obvious to have employed the monitoring
methods of Schumacher to gauge the effectiveness of the medicament. Thus, based on
the reasons of record, the instant claims are deemed unpatentable over the cited art.

#### Conclusion

Claims 12-14 are not allowed.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

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Any inquiry concerning this communication or earlier communications from the examiner should be directed to SAHAR JAVANMARD whose telephone number is (571) 270-3280. The examiner can normally be reached on 8 AM-5 PM MON-FRI (EST).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Sreeni Padmanabhan can be reached on (571) 272-0629. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

/S. J./

Examiner, Art Unit 1617

/SREENI PADMANABHAN/

Supervisory Patent Examiner, Art Unit 1627